



(11) **EP 1 377 690 B1**

(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention
of the grant of the patent:
09.01.2008 Bulletin 2008/02

(51) Int Cl.:
C22B 9/18 ^(2006.01) **C22B 9/14** ^(2006.01)
C22C 19/03 ^(2006.01)

(21) Application number: **02707863.3**

(86) International application number:
PCT/US2002/005510

(22) Date of filing: **25.02.2002**

(87) International publication number:
WO 2002/072897 (19.09.2002 Gazette 2002/38)

(54) **METHOD FOR PRODUCING LARGE DIAMETER INGOTS OF NICKEL BASE ALLOYS**

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PROCEDE DE FABRICATION DE LINGOTS DE GRAND DIAMETRE EN ALLIAGES A BASE DE
NICKEL

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**
Designated Extension States:
SI

(30) Priority: **08.03.2001 US 802064**

(43) Date of publication of application:
07.01.2004 Bulletin 2004/02

(60) Divisional application:
07075914.7

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- **MOYER ET AL.:** 'Advances in triple melting superalloys 718, 706 and 720. The minerals' **METALS & MATERIALS SOCIETY 1994, pages 39 - 48, XP002950951**
- **CORDY ET AL.:** 'Chemistry and structure control in remelted superalloy ingots' **THE PROCEEDINGS VAC. METALL. CONF. SPEC. METALL. MELTING PROCESS 1984, pages 69 - 74, XP002950952**
- **KISSINGER:** 'Trends and near term requirements for GE aircraft engines titanium and nickel base disk alloys' **CONFERENCE OF THE ELECTRON BEAM MELTING AND REFINING - STATE OF THE ART 1991 October 1992, RENO, NEVADA, pages 31 - 40, XP002950953**
- **CHOUDHURY:** 'State of the art of superalloy production for aerospace and other application using VIM/VAR or VIM/ESR' **ISJ INTERNATIONAL vol. 32, no. 5, May 1992, pages 563 - 574, XP000600856**

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Description

[0001] The present invention relates to an improved method for producing large diameter, premium quality ingots of nickel base superalloys. The present invention more particularly relates to a method for producing ingots of nickel base superalloys, including Alloy 718 (UNS N07718) and other nickel base superalloys experiencing significant segregation during casting, and wherein the ingots have a diameter greater than 30 inches (762 mm) and are substantially free of negative segregation, are free of freckles, and are free of other positive segregation. The present invention also is directed to ingots of Alloy 718 having diameters greater than 30 inches (762 mm), as well as to any ingots, regardless of diameter, formed using the method of the invention. The method of the present invention may be applied in, for example, the manufacture of large diameter, premium quality ingots of nickel base superalloys that are fabricated into rotating parts for power generation. Such parts include, for example, wheels and spacers for land-based turbines and rotating components for aeronautical turbines.

DESCRIPTION OF THE INVENTION BACKGROUND

[0002] In certain critical applications, components must be manufactured from nickel base superalloys in the form of large diameter ingots that lack significant segregation. Such ingots must be substantially free of positive and negative segregation, and should be completely free of the manifestation of positive segregation known as "freckles". Freckles are the most common manifestation of positive segregation and are dark etching regions enriched in solute elements. Freckles result from the flow of solute-rich interdendritic liquid in the mushy zone of the ingot during solidification. Freckles in Alloy 718, for example, are enriched in niobium compared to the matrix, have a high density of carbides, and usually contain Laves phase. "White spots" are the major type of negative segregation. These light etching regions, which are depleted in hardener solute elements, such as niobium, typically are classified into dendritic, discrete, and solidification white spots. While there can be some tolerance for dendritic and solidification white spots, discrete white spots are of major concern because they frequently are associated with a cluster of oxides and nitrides that can act as a crack initiator.

[0003] Ingots substantially lacking positive and negative segregation and that are also free of freckles are referred to herein as "premium quality" ingots. Premium quality nickel base superalloy ingots are required in certain critical applications including, for example, rotating components in aeronautical or land-based power generation turbines and in other applications in which segregation-related metallurgical defects may result in catastrophic failure of the component. As used herein, an ingot "substantially lacks" positive and negative segregation when such types of segregation are wholly absent or are present only to an extent that does not make the ingot unsuitable for use in critical applications, such as use for fabrication into rotating components for aeronautical and land-based turbine applications.

[0004] Nickel base superalloys subject to significant positive and negative segregation during casting include, for example, Alloy 718 and Alloy 706. The compositions of Alloys 718 and 706 are well known in the art. The compositions are defined as being:

[0005] Alloy 718 (weight percentages): aluminium 0.20 - 0.8; boron max. 0.006; carbon max. 0.08; cobalt max. 1.00; chromium 17 - 21; copper max. 0.3; manganese max. 0.35; molybdenum 2.8 - 3.3; Nb + Ta 4.75 - 5.5; nickel 50 - 55; phosphorus max. 0.015; sulphur max. 0.015; silicon max. 0.35; titanium 0.65 - 1.15; balance iron and incidental impurities.

[0006] Alloy 706 (weight percentages): aluminium max. 0.40; boron max. 0.006; carbon max. 0.06; cobalt max. 1.00; chromium 14.5 - 17.5; copper max. 0.3; manganese max. 0.35; Nb + Ta 2.5 - 3.3; nickel + cobalt 39.0 - 44.0; phosphorus max. 0.020; sulphur max. 0.015; silicon max. 0.35; titanium 1.5 - 2.0; balance iron and incidental impurities.

[0007] In order to minimize segregation when casting these alloys for use in critical applications, and also to better ensure that the cast alloy is free of deleterious non-metallic inclusions, the molten metallic material is appropriately refined before being finally cast Alloy 718, as well as certain other segregation-prone nickel base superalloys such as Alloy 706 (UNS N09706), are typically refined by a "triple melt" technique which combines, sequentially, vacuum induction melting (VIM), electroslag remelting (ESR), and vacuum arc remelting (VAR). Premium quality ingots of these segregation-prone materials, however, are difficult to produce in large diameters by VAR melting, the last step in the triple melt sequence. In some cases, large diameter ingots are fabricated into single components, so areas of unacceptable segregation in VAR-cast ingots cannot be selectively removed prior to component fabrication. Consequently, the entire ingot or a portion of the ingot may need to be scrapped.

[0008] "Advances in Triple Melting Superalloys 718, 706, and 720" by Moyer et al in "The Minerals, Metals and Materials Society, 1994", pages 39-48, describes that Triple Melting by VIM+ESR+VAR has been highly successful for Alloy 718, large diameter ingots of Alloy 706, and high hardener alloys such as Alloy 720. It further describes how a major advantage of triple melting Alloy 706 is that ingots of up to 36 inches (914mm) diameter can be melted without freckles or positive segregation problems.

[0009] VAR ingots of Alloy 718, Alloy 706, and other nickel base superalloys such as Alloy 600, Alloy 625, Alloy 720, and Waspaloy, are increasingly required in larger weights, and correspondingly larger diameters, for emerging applications. Such applications include, for example, rotating components for larger land-based and aeronautical turbines under

development. Larger ingots are needed not only to achieve the final component weight economically, but also to facilitate sufficient thermomechanical working to adequately break down the ingot structure and achieve all of the final mechanical and structural requirements.

[0010] The melting of large superalloy ingots accentuates a number of basic metallurgical and processing related issues. Heat extraction during melting becomes more difficult with increasing ingot diameter, resulting in longer solidification times and deeper molten pools. This increases the tendency towards positive and negative segregation. Larger ingots and electrodes can also generate higher thermal stresses during heating and cooling. While ingots of the size contemplated by this invention have been successfully produced in several nickel base alloys (for example, Alloys 600, 625, 706, and Waspaloy) Alloy 718 is particularly prone to these problems. To allow for the production of large diameter VAR ingots of acceptable metallurgical quality from Alloy 718 and certain other segregation-prone nickel base superalloys, specialized melting and heat treatment sequences have been developed. Despite these efforts, the largest commercially available premium quality VAR ingots of Alloy 718, for example, are currently 20 inches (508 mm) in diameter, with limited material produced at up to 28-inch (711 mm) diameters. Attempts at casting larger diameter VAR ingots of Alloy 718 material have been unsuccessful due to the occurrence of thermal cracking and undesirable segregation. Due to length restrictions, 28-inch VAR ingots of Alloy 718 weigh no more than about 21,500 lbs (9772 kg). Thus, Alloy 718 VAR ingots in the largest commercially available diameters fall far short of the weights needed in emerging applications requiring premium quality nickel base superalloy material.

[0011] Accordingly, there is a need for an improved method of producing premium quality, large diameter VAR ingots of Alloy 718. There also is a need for an improved method of producing ingots of other segregation-prone nickel base superalloys that are substantially free of negative segregation, are free of freckles, and substantially lack other positive segregation.

BRIEF SUMMARY OF THE INVENTION

[0012] In order to address the above-described needs, the present invention provides a novel method of producing a nickel base superalloy. The method may be used to cast VAR ingots of premium quality from Alloy 718 in diameters greater than 30 inches (762 mm) and having weights in excess of 21,500 lbs (9772 kg). It is believed that the method of the present invention also may be applied in the production of large diameter VAR ingots from other nickel base superalloys subject to significant segregation during casting, such as, for example, Alloy 706.

[0013] The method of the present invention includes the initial step of casting a nickel base superalloy within a casting mold. This may be accomplished by VIM, argon oxygen decarburization (AOD), vacuum oxygen decarburization (VOD), or any other suitable primary melting and casting technique. The cast ingot is subsequently annealed and overaged by heating the alloy at a furnace temperature of at least 1200°F (649°C) for at least 10 hours. (As used herein, "subsequent" and "subsequently" refer to method steps or events that occur immediately one after another, but also refer to method steps or other events that are separated in time and/or by intervening method steps or other events.) In a subsequent step, the ingot is applied as an ESR electrode and is electroslog remelted at a melt rate of at least 8 lbs/min. (3.63 kg/min.). The ESR ingot is transferred to a heating furnace within 4 hours of complete solidification, and is subsequently subjected to a post-ESR heat treatment. The heat treatment includes the steps of holding the alloy at a first furnace temperature of 600°F (316°C) to 1800°F (982°C) for at least 10 hours, and then increasing the furnace temperature, in either a single stage or in multiple stages, from the first furnace temperature to a second furnace temperature of at least 2125°F (1163°C) in a manner that inhibits thermal stresses within the ingot. The ingot is held at the second temperature for at least 10 hours to provide the ingot with a homogenized structure and with minimal Laves phase.

[0014] In some instances, the ESR ingot may be cast with a diameter that is larger than the desired diameter of the VAR electrode to be used in a subsequent step of the method. Therefore, the method of the present invention may include, subsequent to holding the ESR ingot at the second furnace temperature, and prior vacuum arc remelting, mechanically working the ESR ingot at elevated temperature to alter dimensions of the ingot and to provide a VAR electrode of the desired diameter. Thus, after the ESR ingot has been held at the second furnace temperature, it may be further processed in one of several ways, including cooling to a suitable mechanical working temperature or cooling to about room temperature and subsequently reheating to a suitable mechanical working temperature. Alternatively, if adjustment of ingot diameter is unnecessary, the ingot may be directly cooled to room temperature and subsequently processed by vacuum arc remelting without the step of mechanical working. All steps of cooling and reheating the ESR ingot subsequent to holding the ESR ingot at the second temperature are carried out in a manner that inhibits thermal stresses and that will not result in thermal cracking of the ingot.

[0015] In a subsequent step of the present method, the ESR ingot is vacuum arc remelted at a melt rate of 8 to 11 lbs/minute (3.63 to 5 kg/minute) to provide a VAR ingot. The VAR melt rate is preferably 9 to 10.25 lbs/minute (4.09 to 4.66 kg/min), and is more preferably 9.25 to 10.2 lbs/minute (4.20 to 4.63 kg/minute). The VAR ingot preferably has a diameter greater than 30 inches (762 mm), and more preferably has a diameter of at least 36 inches (914 mm).

[0016] The present invention is further directed to a method of producing a nickel base superalloy that is substantially

free of positive and negative segregation and that includes the step of casting in a casting mold an alloy selected from Alloy 718 and other nickel base superalloys subject to significant segregation during casting. The cast ingot is subsequently annealed and overaged by heating at a furnace temperature of at least 1550°F (843°C) for at least 10 hours. The annealed ingot is subsequently electroslog remelted at a melt rate of at least about 10 lbs/min. (4.54 kg/min.), and the ESR ingot is then transferred to a heating furnace within 4 hours of complete solidification. In subsequent steps, the ESR ingot is subjected to a multi-stage post-ESR heat treatment by holding the ingot at a first furnace temperature of 900°F (482°C) to 1800°F (982°C) for at least 10 hours. The furnace temperature is subsequently increased by no more than 100°F/hour (55.6°C/hour) to an intermediate furnace temperature, and is subsequently further increased by no more than 200°F/hour (111°C/hour) to a second furnace temperature of at least 2125°F (1163°C). The ingot is held at the second furnace temperature for at least 10 hours. The ESR ingot may be converted to a VAR electrode of appropriate dimensions, if necessary, and is subsequently vacuum arc remelted at a melt rate of 8 to 11 lbs/minute (3.63 to 5 kg/minute) to provide a VAR ingot. If desired, the VAR ingot may be further processed, such as by a homogenization and/or suitable mechanical conversion to desired dimensions. Representative articles of manufacture that may be fabricated from the ingots fabricated according to the present invention include, for example, wheels and spacers for use in land-based turbines and rotating components for use in aeronautical turbines.

[0017] The reader will appreciate the foregoing details and advantages of the present invention, as well as others, upon consideration of the following detailed description of embodiments of the invention. The reader also may comprehend such additional advantages and details of the present invention upon carrying out or using the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

[0018] The features and advantages of the present invention may be better understood by reference to the accompanying drawings in which:

Figure 1 is a diagram generally illustrating of one embodiment of the method of the present invention, wherein the ESR ingot has a 40-inch diameter and is converted to a 32-inch diameter VAR electrode prior to vacuum arc remelting; Figure 2 is a diagram generally illustrating a second embodiment of the method of the present invention, wherein the ESR ingot has a 36-inch diameter and is converted to a 32-inch diameter VAR electrode prior to vacuum arc remelting ; and

Figure 3 is a diagram of a third embodiment of the method of the present invention, wherein a 33-inch diameter ESR ingot is cast and is suitable without mechanical conversion for use as the VAR electrode.

DETAILED DESCRIPTION OF EMBODIMENTS OF THE INVENTION

[0019] The method of the present invention allows for the production of premium quality, large diameter ingots from Alloy 718, a nickel base superalloy that is prone to segregation on casting. Previous to the development of the present method, the heaviest commercially available ingots of Alloy 718 were limited to about 28 inches (711 mm) in diameter, with maximum weights of about 21,500 lbs (9773 kg) because of length/diameter limitations. The inventors have successfully produced premium quality ingots of Alloy 718 with diameters greater than 30 inches (762 mm) and at least 36 inches (914 mm) by the present method. These ingots weighed as much as 36,000 lbs (16,363 kg), well in excess of the previous maximum weight for premium quality 718 Alloy VAR ingots. The inventors believe that the method of the present invention may be used to produce VAR ingots of other nickel base superalloys that typically experience significant segregation during casting. Such other alloys include, for example, Alloy 706.

[0020] The method of the present invention includes the step of casting a nickel base superalloy within a casting mold. As noted, the nickel base alloy may be, for example, Alloy 718. Alloy 718 has the following broad composition, all in weight percentages: about 50.0 to about 55.0 nickel; about 17 to about 21.0 chromium; 0 up to about 0.08 carbon; 0 up to about 0.35 manganese; 0 up to about 0.35 silicon; about 2.8 up to about 3.3 molybdenum; at least one of niobium and tantalum, wherein the sum of niobium and tantalum is about 4.75 up to about 5.5; about 0.65 up to about 1.15 titanium; about 0.20 up to about 0.8 aluminum; 0 up to about 0.006 boron; and iron and incidental impurities. Alloy 718 is available under the trademark Allvac 718 from the Allvac division of Allegheny Technologies Incorporated, Pittsburgh, Pennsylvania. Allvac 718 has the following nominal composition (in weight percentages) when cast in larger VAR ingot diameters: 54.0 nickel; 0.5 aluminum; 0.01 carbon; 5.0 niobium; 18.0 chromium; 3.0 molybdenum; 0.9 titanium; and iron and incidental impurities.

[0021] Any suitable technique may be used to melt and cast the alloy within a casting mold. Suitable techniques include, for example, VIM, AOD, and VOD. The choice of melting and casting technique is often dictated by a combination of cost and technical issues. Electric arc furnace/AOD melting facilitates the use of low cost raw materials, but tends to be lower in yield than VIM melting, particularly if bottom pouring is used. As the cost of raw materials increases, the higher yield from VIM melting may make this a more economical approach. Alloys containing higher levels of reactive

elements may require VIM melting to ensure adequate recovery. The need for low gaseous residual contents, particularly nitrogen, also may dictate the use of VIM melting to reach the desired levels.

[0022] After the alloy has been cast, it may be held within the mold for a certain period to ensure sufficient solidification so that it may be stripped safely from the casting mold. Those of ordinary skill in the art may readily determine a sufficient time, if any, to hold the cast ingot within mold. That time will depend on, for example, the size and dimensions of the ingot, the parameters of the casting operation, and the composition of the ingot.

[0023] Subsequent to removing the cast ingot from casting mold, it is placed in a heating furnace and is annealed and overaged by heating at a furnace temperature of least 1200°F (649°C) for at least 10 hours. Preferably, the ingot is heated at a furnace temperature of at least 1200°F (649°C) for at least 18 hours. A more preferable heating temperature is at least 1550°F (843°C). The annealing and overaging heat treatment is intended to remove residual stresses within the ingot created during solidification. As ingot diameter increases, residual stresses become more of a concern because of increased thermal gradients within the ingot and the degree of microsegregation and macrosegregation increases, raising the sensitivity to thermal cracking. When residual stresses become excessive, thermal cracks can initiate. Some thermal cracks may be catastrophic, resulting in the need to scrap the product. Cracking may also be more subtle and result in melting irregularities and subsequent unacceptable segregation. One type of melting irregularity known as a "melt rate cycle" is caused by thermal cracks introduced into the ESR and VAR electrode that interrupt heat conduction along the electrode from the tip that is melting. This concentrates the heat below the crack, which causes the melt rate to increase as the melting interface approaches the crack. When the crack is reached, the end of the electrode is relatively cold, making the melting process suddenly slower. As the crack region melts, the melt rate gradually increases until a steady state temperature gradient is reestablished in the electrode and the nominal melt rate is reached.

[0024] In a subsequent step, the ingot is used as an ESR electrode to form an ESR ingot. The inventors have determined that an ESR melt rate of at least about 8 lbs/minute (3.63 kg/minute), and more preferably at least 10 lbs/minute (4.54 kg/minute) should be used to provide an ESR ingot suitable for further processing to a large diameter VAR ingot. Any suitable flux and flux feed rate may be used, and those having ordinary skill in the art may readily determine suitable fluxes and feed rates for a given ESR process. To some extent, the suitable melting rate will depend on the desired ESR ingot diameter and should be selected to provide an ESR ingot of a solid construction (*i.e.*, substantially lacking voids and cracks), having reasonably good surface quality, and lacking excessive residual stresses to inhibit thermal cracking. The general operation of ESR equipment and the general manner of conducting the remelting operation are well known to those of ordinary skill in the art. Such persons may readily electroslag remelt an ESR electrode of a nickel base superalloy, such as Alloy 718, at the melt rate specified in the present method without further instruction.

[0025] Once the electroslag remelting operation has been completed, the ESR ingot may be allowed to cool in the crucible to better ensure that all molten metal has solidified. The minimum suitable cool time will largely depend on ingot diameter. Once removed from the crucible, the ingot is transferred to a heating furnace so that it may be subjected to a novel post-ESR heat treatment according to the present invention and as follows.

[0026] The inventors have discovered that in the production of large diameter ingots of Alloy 718, it is important that the ESR ingot is hot transferred into the heating furnace and that the post-ESR heat treatment be initiated within 4 hours from the complete solidification of the ESR ingot. Once the ESR ingot has been transferred to the heating furnace, the post-ESR heat treatment is initiated by holding the ingot at a first furnace temperature in the range of at least 600°F (316°C) up to 1800°F (982°C) for at least 10 hours. More preferably, the furnace temperature range is least 900°F (482°C) up to 1800°F (982°C). It also is preferred that the heating time at the selected furnace temperature is at least 20 hours.

[0027] After the step of holding the furnace temperature for at least 10 hours, the heating furnace temperature is increased from the first furnace temperature up to a second furnace temperature of at least 2125°F (1163°C), and preferably at least 2175°F (1191°C), in a manner that inhibits the generation of thermal stresses within the ESR ingot. The increase in furnace temperature up to the second furnace temperature may be performed in a single stage or as a multiple-stage operation including two or more heating stages. The inventors have determined that a particularly satisfactory sequence of increasing temperature from the first to the second furnace temperatures is a two-stage sequence including: increasing furnace temperature from the first temperature by no greater than 100°F/hour (55.6°C/hour), and preferably about 25°F/hour (13.9°C/hour), to an intermediate temperature; and then further increasing furnace temperature from the intermediate temperature by no greater than 200°F/hour (111°C/hour), and preferably about 50°F/hour (27.8°C/hour), to the second furnace temperature. Preferably, the intermediate temperature is at least 1000°F (538°C), and more preferably is at least 1400°F (760°C).

[0028] The ESR ingot is held at the second furnace temperature for at least 10 hours. The inventors have determined that after being held at the second furnace temperature, the ingot should exhibit a homogenized structure and include only minimal Laves phase. In order to better ensure that that desired structure and the desired degree of annealing is achieved, the ESR ingot is preferably held at the second furnace temperature for at least 24 hours, and is more preferably held at the second furnace temperature for about 32 hours.

[0029] After the ESR ingot has been held at the second furnace temperature for the specified period, it may be further

processed in one of several ways. If the ESR ingot will not be mechanically worked, it may be cooled from the second furnace temperature to room temperature in a manner that inhibits thermal cracking. If the ESR ingot has a diameter that is greater than the desired diameter of the VAR electrode, the ESR ingot may be mechanically worked such as by, for example, hot forging. The ESR ingot may be cooled from the second furnace temperature to a suitable mechanical working temperature in a manner selected to inhibit thermal cracking. If, however, the ESR ingot has been cooled below a suitable working temperature, it may be reheated to the working temperature in a fashion that inhibits thermal cracking and may then be worked to the desired dimensions.

[0030] The inventors have determined that when cooling the ESR ingot from the second furnace temperature, it is desirable to do so in a controlled manner by reducing furnace temperature from the second furnace temperature while the ingot remains in the heating furnace. A preferred cooling sequence that has been shown to prevent thermal cracking includes: reducing the furnace temperature from the second furnace temperature at a rate no greater than 200°F/hour (111°C/hour), and preferably at about 100°F/hour (55.6°C/hour), to a first intermediate temperature not greater than 1750°F (954°C), and preferably not greater than 1600°F (871 °C); holding at the first intermediate temperature for at least 10 hours, and preferably at least 18 hours; further reducing the furnace temperature from the first intermediate temperature at a rate not greater than 150°F/hour (83.3°C/hour), and preferably about 75°F/hour (41.7°C/hour), to a second intermediate temperature not greater than 1400°F (760°C), and preferably not greater than 1150°F (621 °C); holding at the second intermediate temperature for at least 5 hours, and preferably at least 7 hours; and subsequently air cooling the ingot to room temperature. Once cooled to room temperature, the ingot should exhibit an overaged structure of delta phase precipitates.

[0031] If the ESR ingot is cooled from the second furnace temperature to a temperature at which mechanical working will be carried out, then the relevant portion of the cooling sequence just described may be used to achieve the working temperature. For example, if the ESR ingot is being heated in a heating furnace at a second furnace temperature of 2175°F (1191°C) and is to be hot forged at a forging temperature of 2025°F (1107°C), the ESR ingot may be cooled by reducing the furnace temperature from the second furnace temperature at a rate no greater than 200°F/hour (111°C/hour), and preferably at about 100°F/hour, to the forging temperature.

[0032] The inventors have determined that if the ESR ingot has been cooled from the second furnace temperature to a temperature at or near room temperature, then heating the ingot back to a suitable mechanical working temperature may be conducted using the following sequence in order to inhibit thermal cracking: charge the ingot to a heating furnace and heat the ingot at a furnace temperature less than 1000°F (556°C) for at least 2 hours; increase the furnace temperature at less than 40°F/hour (22.2°C/hour) to less than 1500°F (816°C); further increase the furnace temperature at less than 50°F/hour (27.8°C/hour) to a suitable hot working temperature less than 2100°F (1149°C); and hold the ingot at the working temperature for at least 4 hours. In an alternate heating sequence developed by the inventors, the ESR ingot is placed in a heating furnace and the following heating sequence is followed: the ingot is heated at a furnace temperature of at least 500°F (260°C), and preferably at 500-1000°F (277-556°C), for at least 2 hours; the furnace temperature is increased by about 20-40°F/hour (11.1-22.2°C/hour) to at least 800°F (427°C); the furnace temperature is further increased by about 30-50°F/hour (16.7-27.8°C/hour) to at least 1200°F (649°C); the furnace temperature is further increased by about 40-60°F/hour (22.2-33.3°C/hour) to a hot working temperature less than 2100°F (1149°C); and the ingot is held at the hot working temperature until the ingot achieves a substantially uniform temperature throughout.

[0033] If the ESR ingot has been cooled or heated to a desired mechanical working temperature, it is then worked in any suitable manner, such as by press forging, to provide a VAR electrode having a predetermined diameter. Reductions in diameter may be necessitated by, for example, limitations on available equipment. As an example, it may be necessary to mechanically work an ESR ingot having a diameter of about 34 to about 40 inches (about 864 to about 1016 mm) to a diameter of 34 inches (about 864 mm) or less so that it may suitably be used as the VAR electrode on available VAR equipment.

[0034] To this point, the ESR ingot will have been subjected to the post-ESR heat treatment. It also has assumed, either as cast on the ESR apparatus or after mechanical working, a suitable diameter for use as the VAR electrode. The ESR ingot may then be conditioned and cropped to adjust its shape to that suitable for use as a VAR electrode, as is known in the art. The VAR electrode is subsequently vacuum arc remelted at a rate of 8 to 11 lbs/minute (3.63 to 5 kg/minute) in a manner known to those of ordinary skill in the art to provide a VAR ingot of the desired diameter. The VAR melt rate is preferably 9 to 10.25 lbs/minute (4.09 to 4.66 kg/min), and is even more preferably 9.25 to 10.2 lbs/minute (4.20 to 4.63 kg/minute). The inventors have determined that the VAR melt rate is critical to achieving premium quality VAR ingots of Alloy 718 material.

[0035] The cast VAR ingot may be further processed, if desired. For example, the VAR ingot may be homogenized and overaged using techniques conventional in the production of commercially available larger diameter nickel base superalloy VAR ingots.

[0036] Nickel base superalloy ingots produced by the method of the present invention may be fabricated into articles of manufacture by known manufacturing techniques. Such articles would naturally include certain rotating components adapted for use in aeronautical and land-based power generation turbines.

[0037] Examples of the method of the present invention follow.

Example 1

[0038] Figure 1 is a diagram generally depicting an embodiment of the method of the present invention adapted for producing premium quality ingots of Alloy 718 with diameters greater than 30 inches. It will be apparent that the embodiment of the present method shown in Figure 1 is, in general, a triple-melt process including steps of VIM, ESR, and VAR. As indicated in Figure 1, a heat of Alloy 718 was prepared by VIM and cast to a 36-inch diameter VIM electrode suitable for use as an ESR electrode in a subsequent step. The VIM ingot was allowed to remain in the casting mold for 6 to 8 hours after casting. The ingot was then stripped from the mold and transferred hot to a furnace, where it was annealed and overaged at 1550°F (843°F) for 18 hours minimum.

[0039] After the anneal/overage step, the ingot surface was ground to remove scale. The ingot was then transferred hot to an ESR apparatus, where it was used as the ESR consumable electrode and was electroslag remelted to form a 40-inch ESR ingot. As is well known, an ESR apparatus includes an electric power supply that is in electrical contact with the consumable electrode. The electrode is in contact with a slag disposed in a water-cooled vessel, typically constructed of copper. The electric power supply, which is typically AC, provides a high amperage, low voltage current to a circuit that includes the electrode, the slag, and the vessel. As current passes through the circuit, electrical resistance heating of the slag increases its temperature to a level sufficient to melt the end of the electrode in contact with the slag. As the electrode begins to melt, droplets of molten material form, and an electrode feed mechanism advances the electrode into the slag to provide the desired melt rate. The molten material droplets pass through the heated slag, which removes oxide inclusions and other impurities. Determining the proper melt rate is crucial to provide an ingot that is substantially homogenous and free of voids, and that has a reasonably good quality surface. Here, the inventors determined through experimentation that a melt rate of 14 lbs/min. provided a suitably homogenous and defect-free ESR ingot.

[0040] After the 40-inch ESR ingot was cast, it was allowed to cool within the mold for 2 hours and then subjected to the following post-ESR heat treatment. The heat treatment prevented thermal cracking in the ingot in subsequent processing. The ESR ingot was removed from the mold and hot transferred to a heating furnace where it was maintained at about 900°F (482°C) for 20 hours. Furnace temperature was then increased by about 25°F/hour (13.9°C/hour) to about 1400°F (760°C). Furnace temperature was then further increased at a rate of about 50°F/hour (27.8°C/hour) to about 2175°F (1191 °C), and the ingot was held at 2175°F (1191 °C) for at least 32 hours. The ingot was then cooled by reducing furnace temperature about 100°F/hour (55.6°C/hour) to about 1600°F (871°C). That temperature was maintained for at least 18 hours. The ingot was then further cooled by reducing the furnace temperature about 75°F/hour (41.7°C/hour) to about 1150°F, and the temperature was held there for about 7 hours. The ingot was removed from the furnace and allowed to air cool.

[0041] The 40-inch diameter of the ESR ingot was too large to be vacuum arc remelted using the available VAR apparatus. Therefore, the ingot was press forged to a 32-inch diameter suitable for use on the VAR apparatus. Before forging, the ingot was heated in a furnace to a suitable press forging temperature by a heating sequence developed by the present inventors to prevent thermal cracking. The ingot was first heated at 500°F (260°C) for 2 hours. Furnace temperature was then ramped up at 20°F/hour (11.1°C/hour) to 800°F (427°C), increased by 30°F/hour (16.7°C/hour) to 1200°F (649°C), and then further increased by 40°F/hour (22.2°C/hour) to 2025°F (1107°C), where it was maintained for about 8 hours. The ingot was then press forged to a 32-inch diameter, reheating to forging temperature as needed. The 32-inch VAR electrode was maintained at about 1600°F (871 °C) for a minimum of 20 hours and then conditioned and bandsaw cropped to flatten its ends.

[0042] The inventors have discovered that only a narrow and specific VAR melting range will produce a substantially segregation-free VAR ingot, and that VAR control is especially critical during start-up to avoid macrosegregation. The 32-inch VAR electrode was vacuum arc remelted to a 36-inch VAR ingot at a melt rate of about 9.75 lbs/min., which must be controlled within a narrow window. The VAR ingot was then homogenized using a standard furnace homogenization heating cycle, and was then overaged at 1600°F (871 °C) for 20 hours minimum.

[0043] The weight of the 36-inch VAR ingot was significantly in excess of the 21,500 lb (9772 kg) weight of commercially available 28-inch diameter Alloy 718 ingots. Product from the 36-inch ingot was ultrasonically and macro slice inspected, and was found to be free of freckles, and was substantially free of cracks, voids, negative segregation, and other positive segregation. The ESR ingot was considered to be premium quality and suitable for fabrication into parts used in critical applications, such as rotating parts for land-based and aeronautical power generation turbines.

Example 2

[0044] In the above example, the ESR ingot had a diameter in excess of that which could be used on the available VAR apparatus, which accommodated a VAR electrode of up to about 34 inches (863 mm). This necessitated that the diameter of the ESR ingot be adjusted by mechanical working. This, in turn, required that the inventors develop a suitable

ESR ingot heating sequence to heat the ESR ingot to forging temperature while preventing the occurrence of thermal cracking during forging. If the diameter of the ESR ingot were to more closely approximate the maximum diameter usable on the available VAR apparatus, then the ESR ingot would be less prone to thermal cracking. Press forging or other mechanical working of the ESR ingot may be wholly unnecessary if the size of the ESR ingot were suitable for use directly on the available VAR apparatus. In such case, the ESR ingot could be delivered to the VAR apparatus immediately after the post-ESR heat treatment steps.

[0045] Figure 2 is a diagram generally depicting a prophetic embodiment of a triple-melt process according to the present invention wherein the ESR apparatus may be used to cast a 36-inch ESR ingot. Because the ESR ingot has a diameter that is less than the 40-inch diameter of the ESR ingot cast in Example 1, there would be less risk of ingot cracking or other working-induced imperfections. In addition, the reduced diameter and greater length of the ESR ingot would reduce the likelihood that the ESR ingot would crack or suffer from significant segregation once cast.

[0046] As indicated in Figure 2, the VIM electrode is cast to a 33-inch diameter ingot. The VIM ingot is then hot transferred and may be annealed and overaged as described in Example 1. In particular, the VIM ingot is allowed to remain in the casting mold for 6 to 8 hours before being stripped and loaded into the heat-treating furnace. It is believed that the hold time in the casting mold could be reduced for smaller diameter VIM ingots. The 33-inch VIM ingot is then electroslog remelted by the process generally described in Example 1. The ingot is then hot transferred and subjected to a post-ESR heat treatment as described above in Example 1. Subsequent to the post-ESR heat treatment, the ESR ingot is ramped up to forging temperature and press forged to 32-inch diameter as generally described in Example 1. The 32-inch forging is overaged and then vacuum arc remelted to a 36-inch VAR ingot as generally described in Example 1. The VAR ingot may then be homogenized by standard homogenization treatments, or may be suitably processed in other ways. It is believed that a premium quality Alloy 718 VAR ingot, comparable to the ingot produced by the method of Example 1, would result.

Example 3

[0047] Figure 3 is a diagram an alternative prophetic embodiment of a triple-melt process within the present invention wherein the 30-inch diameter of the as-cast ESR ingot is directly suitable for use with the ESR apparatus. A 30-inch VIM electrode is electroslog remelted to a 33-inch ESR ingot. The ESR ingot is hot transferred and heat treated as described in Example 1, and is then vacuum arc remelted, without reduction in diameter, to a 36-inch diameter VAR ingot. The VAR ingot may then be homogenized and further processed as described in Example 1. The process depicted in Figure 3 differs from that of Figure 1 only in that the diameters of the VIM electrode and ESR ingot differ from those of Example 1, and no press forging operation or ramped heat-up to forging temperature are needed. A premium quality 36-inch diameter Alloy 718 ingot would result.

Example 4

[0048] Several VAR ingots of Allvac 718 material having diameters greater than 30 inches were prepared by the method of the present invention and inspected. Parameters of the several runs are set forth in the following chart. In several of the runs, various VAR melt rates were evaluated to determine the effects on quality of the resulting VAR ingot.

Step	Heat 215G	Heat 420G	Heat 533G	Heat 631 G	Heat 729G
VIM Electrode Diameter	36	36	36	36	36
VIM Anneal/Overage	1550°F (843°C) for 13 hours 24 minutes	1550°F (843°C) for 16 hours 48 minutes	1550°F (843°C) for 15 hours 55 minutes	1550°F (843°C) for 41 hours	1550°F (843°C) for 29 hours
Flux	60F-20-0-20 + TiO ₂	60F-20-0-20 + TiO ₂	60F-20-0-20 + TiO ₂	60F-20-0-20 + TiO ₂	60F-20-0-20 + TiO ₂
ESR Melt Rate	14 lbs/minute	14 lbs/minute	14 lbs/minute	14 lbs/minute	14 lbs/minute
Crucible Cool Time	1.5 hours (1 hour 50 minutes total transfer time)	2 hours	2 hours	2 hours (+ 20 minutes to strip to hot box)	2 hours (+ 30 minutes to strip to hot box)
ESR Ingot Diameter	40 inches	40 inches	40 inches	40 inches	40 inches

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(continued)

Step	Heat 215G	Heat 420G	Heat 533G	Heat 631 G	Heat 729G
Post ESR Heat Treatment	900°F (482°C) for 33 hours 22 minutes. 1150°F (621°C) for 7 hours. Ramp up at 25°F/hour (13.8°C/hour) to 1300°F (704°C), then 50°F/hour (27.7°C/hour) to 1650°F (899°C), and 75°F/hour (41.6°C/hour) to 2175°F (1191°C). Hold for 24 hours at 2175°F (1191°C). Lower to 2025°F (1107°C), hold for 6 hours and forge.	900°F (482°C) for 28 hours. 1150°F (621°C) for 19 hours. Ramp up at 25°F/hour (13.8°C/hour) to 1300°F (704°C), then 50°F/hour (27.7°C/hour) to 1650°F (899°C), and 75°F/hour (41.6°C/hour) to 2175°F (1191°C). Hold for 24 hours at 2175°F (1191°C). Lower to 2025°F (1107°C), hold for 9 hours and forge.	900°F (482°C) for 21 hours. 1150°F (621°C) for 4 hours. Ramp up at 25°F/hour (13.8°C/hour) to 1300°F (704°C), then 50°F/hour (27.7°C/hour) to 1650°F (899°C), and 75°F/hour (41.6°C/hour) to 2175°F (1191°C). Hold for 24 hours at 2175°F (1191°C). Lower to 2025°F (1107°C), hold for 69.5 hours and forge.	900°F (482°C) for 33 hours. 1150°F (621°C) for 4 hours. Ramp up at 25°F/hour (13.8°C/hour) to 1300°F (704°C), then 50°F/hour (27.7°C/hour) to 1650°F (899°C), and 75°F/hour (41.6°C/hour) to 2175°F (1191°C). Hold for 24 hours at 2175°F (1191°C). Air cool.	900°F (482°C) for 42.5 hours. Ramp up at 25°F/hour (13.8°C/hour) to 1400°F (760°C), then 50°F/hour (27.7°C/hour) to 2175°F (1191°C). Hold for 32 hours at 2175°F (1191°C). Ramp furnace down at 100°F/hour (55.5°C/hour) to 1600°F (871°C) and hold for 18 hours min. Ramp down at 75°F/hour (41.6°C/hour) to 1150°F (621°C) and hold for 7 hours min. Air cool.
Press	Forge to 31-15/16 inches in three operations	Forge to 31-15/16 inches in three operations	Forge to 31-15/16 inches in five operations	Reheat at 500°F (260°C) for 8 hours, ramp at 25°F/hour (13.8°C/hour) to 1300°F (704°C). Ramp at 50°F/hour (27.7°C/hour) to 2025°F (1107°C). Hold at 2025°F (1107°C) and forge	Reheat at 500°F (260°C) for 3.5 hours, ramp at 20°/hour (11.1°C/hour) to 800°F (427°C), ramp at 30°F/hour (16.7°C/hour) to 1200°F (649°C), ramp at 40°F/hour to 2025°F (1107°C). Hold 16 hours at 2025°F (1107°C) and press, reheating as needed.
Forgeback Diameter	31-15/16 inches	31-15/16 inches	31-15/16 inches	Not applicable	32 inches
Overage	1600°F (871°C) for 21 hours and air cool	1600°F (871°C) for 23.5 hours and air cool	1600°F (871°C) for 25 hours and air cool	Not applicable	1600°F (871°C) for 20 hours and air cool
Melt Rate	3 trialed: 9.75, 10.5, and 9.0 lbs/minute	2 trialed: 10.0 and 9.5 lbs/minute	3 trialed: 10.2, 9.25, and 9.75 lbs/minute	Not applicable	9.75

(continued)

Step	Heat 215G	Heat 420G	Heat 533G	Heat 631 G	Heat 729G
VAR Ingot Diameter/Weight	36 inches 27,355 pounds	36 inches 28,570 pounds	36 inches 30,744 pounds	Not applicable	36 inches 37,880 pounds
Homogenize	Yes	Yes	Yes	Not applicable	Yes
Comments	Positive segregation found at highest melt rate. Two ultrasonic indications found in VAR start up area but no freckles found. Material melted under steady state conditions acceptable for premium quality applications.	No ultrasonic indications. Material melted under steady state conditions acceptable for premium quality applications.	No ultrasonic indications. Material melted under steady state conditions acceptable for premium quality applications.	ESR ingot cracked on removal from reheat furnace. Ingot scrapped.	Sound, crack free ingot after VAR

[0049] Evaluation of the VAR ingots was conducted on 25 cm (10-inch) diameter billet produced by draw forging the VAR ingots, followed by GFM forging to final diameter. The forged billets were peeled and polished to remove surface irregularities after which they were ultrasonic inspected for internal cracks and voids that are usually associated with areas of negative segregation. Transverse slices cut from several locations along the length of the billets representing all melt rates were then chemically etched to reveal areas of negative and positive segregation. The absence of sonic indications and segregation defects was sufficient to classify the material as being of premium quality.

[0050] It is to be understood that the present description illustrates those aspects of the invention relevant to a clear understanding of the invention. Certain aspects of the invention that would be apparent to those of ordinary skill in the art and that, therefore, would not facilitate a better understanding of the invention have not been presented in order to simplify the present description. Although the present invention has been described in connection with certain embodiments, those of ordinary skill in the art will, upon considering the foregoing description, recognize that many modifications and variations of the invention may be employed. All such variations and modifications of the invention are intended to be covered by the foregoing description and the following claims.

Claims

1. A method of producing a nickel base superalloy that is substantially free of positive and negative segregation, the method comprising:

casting an alloy that is a nickel base superalloy within a casting mold;
annealing and overaging the alloy by heating the alloy at at least 1200°F (649°C) for at least 10 hours;
electroslag remelting the alloy at a melt rate of at least 8lbs/min (3.63kg/min);
transferring the alloy to a heating furnace within 4 hours of complete solidification;
holding the alloy within the heating furnace at a first temperature of 600°F (316°C) to 1800°F (982°C) for at least 10 hours;
increasing the furnace temperature from the first temperature to a second temperature of at least 2125°F (1163°C) in a manner to inhibit thermal stresses within the alloy;
holding at the second temperature for at least 10 hours;
vacuum arc remelting a VAR electrode of the alloy at a melt rate of 8 to 11 lbs/minute (3.63 to 5kg/minute) to provide a VAR ingot.

2. The method of claim 1, wherein the nickel base alloy is one of Alloy 718 and Alloy 706.

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3. The method of claim 1 or claim 2, wherein casting the nickel base alloy comprises melting and optionally refining the alloy by at least one of vacuum induction melting, argon oxygen decarburization, and vacuum oxygen decarburization.
- 5 4. The method of any one of the preceding claims, wherein annealing and overaging the alloy comprises heating the alloy at at least 1200°F (649°C) for at least 18 hours.
5. The method of any one of the preceding claims, wherein annealing and overaging the alloy comprises heating the alloy at at least 1550°F (843°C) for at least 10 hours.
- 10 6. The method of any one of the preceding claims, wherein electroslag remelting the alloy comprises electroslag remelting at a melt rate of at least 10 lbs/minute (4.54 kg/minute).
7. The method of any one of the preceding claims, wherein holding the alloy within the heating furnace comprises holding the alloy at a furnace temperature of at least 600°F (316°C) up to 1800°F (982°C) for at least 20 hours.
- 15 8. The method of claim 7, wherein holding the alloy within the heating furnace comprises holding the alloy at a furnace temperature of at least 900°F (482°C) up to 1800°F (982°C) for at least 10 hours.
- 20 9. The method of any one of the preceding claims, wherein increasing the furnace temperature comprises increasing the furnace temperature from the first temperature to the second temperature in a multi-stage manner comprising:
 - increasing the furnace temperature from the first temperature by no greater than 1.00°F/hour (55.6°C/hour) to an intermediate temperature; and
 - 25 further increasing the furnace temperature by no greater than 200°F/hour (111°C/hour) from the intermediate temperature to the second temperature.
10. The method of claim 9, wherein the first temperature is less than 1000°F (583°C) and the intermediate temperature is at least 1000°F (583°C).
- 30 11. The method of claim 9, wherein the first temperature is less than 1400°F (760°C) and the intermediate temperature is at least 1400°F (760°C).
12. The method of any one of the preceding claims, wherein the second temperature is at least 21.75°F (1191°C).
- 35 13. The method of any one of the preceding claims, wherein the alloy is held at the second temperature for at least 24 hours.
14. The method of any one of the preceding claims, wherein electroslag remelting the alloy provides an ESR ingot having a diameter that is greater than a desired diameter of the VAR electrode, the method further comprising, subsequent to holding at the second temperature:
 - 40 mechanically working the ESR ingot to alter dimensions of the ingot and to provide a VAR electrode with the desired diameter.
- 45 15. The method of claim 14, wherein the ESR ingot has a diameter of about 34 inches (864 mm) to about 40 inches (1016 mm) and the VAR electrode has a smaller diameter no greater than about 34 inches (864 mm).
16. The method of claim 14 or 15, further comprising, subsequent to holding the alloy at the second temperature and prior to mechanically working the ESR ingot:
 - 50 cooling the alloy to a mechanical working temperature at a cooling rate not greater than 200°F/hour (111°C/hour).
17. The method of any one of the preceding claims, further comprising, subsequent to holding the alloy at the second temperature and prior to vacuum arc remelting the VAR electrode:
 - 55 cooling the alloy from the second temperature to room temperature by a cooling process comprising reducing the furnace temperature at a rate not greater than 200°F/hour (111°C/hour) from the second temperature to a

first intermediate temperature not greater than 1750°F (982°C) and holding at the first intermediate temperature for at least 10 hours.

18. The method of claim 17, wherein cooling the alloy further comprises:

reducing the furnace temperature at a rate not greater than 150°F/hour (83.3°C/hour) from the first intermediate temperature to a second intermediate temperature not greater than 1400°F (760°C) and holding at the second intermediate temperature for at least 5 hours.

19. The method of claim 8, wherein subsequent to holding at the second intermediate temperature, the alloy is cooled in air to about room temperature.

20. The method of any one of the preceding claims, further comprising, subsequent to holding the alloy at the second temperature and prior to mechanically working the ESR ingot:

cooling the alloy from the second temperature to about room temperature in a manner that inhibits thermal stresses in the alloy; and
heating the alloy to a suitable mechanical working temperature in a manner that inhibits thermal stresses in the alloy.

21. The method of claim 20, wherein heating the alloy to a suitable mechanical working temperature comprises:

heating the alloy within a heating furnace at a furnace temperature of at least 500°F (260°C) for at least 2 hours; increasing the furnace temperature by at least about 20°F/hour (11.1°C/hour) to at least 800°F (427°C); further increasing the furnace temperature by at least about 30°F/hour (16.7°C/hour) to at least 1200°F (649°C); and
further increasing the furnace temperature by at least about 40°F/hour (22.2°C/hour) to a temperature of at least 2025°F (1107°C), and holding at the temperature until the alloy achieves a substantially uniform temperature throughout.

22. The method of any one of the preceding claims wherein the method comprises:

casting a nickel base alloy in a casting mold, wherein the nickel base superalloy is Alloy 71.8; annealing and overaging the alloy by heating the alloy at at least 1550°F (843°C) for at least 10 hours; electroslog remelting the alloy at a melt rate of at least 10 lbs/min (4.54 kg/min); transferring the alloy to a heating furnace within 4 hours of complete solidification after electroslog remelting; holding the alloy within the heating furnace at a first furnace temperature of 940°F (482°C) to 1800°F (982°C) for at least 10 hours; increasing the furnace temperature by no greater than 100°F/hour (55.6°C/hour) to an intermediate furnace temperature; and
further increasing the furnace temperature by no greater than 200°F/hour (111°C/hour) from the intermediate furnace temperature to a second furnace temperature of at least 2125°F (1163°C), and holding at the second temperature for at least 10 hours; and
vacuum arc remelting a VAR electrode of the alloy at a melt rate of 9 to 10.25 lbs/minute (4.09 to 4.66 kg/minute) to provide a VAR ingot.

23. The method of any one of the preceding claims, wherein the VAR ingot has a diameter greater than 30 inches (762 mm).

24. The method of any one of the preceding claims, wherein the VAR ingot has a diameter of at least 36 inches (914 mm).

25. The method of any one of the preceding claims, wherein the weight of the VAR ingot is greater than 21,500 lbs (9772 kg).

26. The method of any one of the preceding claims, wherein the nickel base alloy comprises
50.0 to 55.0 weight percent nickel;
17 to 21.0 weight percent chromium;
0 up to 0.08 weight percent carbon;

0 up to 0.35 weight percent manganese;
 0 up to 0.35 weight percent silicon;
 2.8 up to 3.3 weight percent molybdenum;
 at least one of niobium and tantalum wherein the sum of niobium and tantalum is 4.75 up to 5.5 weight percent;
 0.65 up to 1.15 weight percent titanium;
 0.20 up to 0.8 weight percent aluminum;
 0 up to 0.006 weight percent boron; and
 balance iron and incidental impurities.

27. The method of claim 26, wherein the nickel base alloy consists essentially of:

54.0 weight percent nickel;
 0.5 weight percent aluminum;
 0.01 weight percent carbon;
 5.0 weight percent niobium;
 18.0 weight percent chromium;
 3.0 weight percent molybdenum;
 0.9 weight percent titanium; and
 balance iron and incidental impurities.

28. The method of claim 22, wherein electroslog remelting the alloy provides an ESR ingot having a diameter that is greater than a desired diameter of the VAR electrode, the method further comprising:

cooling the alloy from the second temperature to a suitable mechanical working temperature and then mechanically working the alloy to provide a VAR electrode with the desired diameter.

29. The method of claim 22, wherein electroslog remelting the alloy provides an ESR ingot having a diameter that is greater than a desired diameter of the VAR electrode, the method further comprising:

cooling the alloy from the second temperature to about room temperature in a manner that inhibits thermal stresses in the alloy,
 heating the alloy to a suitable mechanical working temperature in a manner that inhibits thermal stresses in the alloy;
 mechanically working the alloy to provide a VAR electrode with the desired diameter.

Patentansprüche

1. Verfahren zur Herstellung einer Superlegierung auf Nickelbasis, die im Wesentlichen frei von positiver und negativer Seigerung ist, wobei das Verfahren Folgendes umfasst:

Gießen einer Legierung, die eine Superlegierung auf Nickelbasis ist, in einer Gießform;
 Glühen und Überaltern der Legierung durch Erwärmen der Legierung bei mindestens 1200°F (649°C) über eine Dauer von mindestens 10 Stunden;
 Elektroschlacke-Umschmelzen der Legierung mit einer Schmelzrate von mindestens 8 lbs/min (3,63 kg/min);
 Verbringen der Legierung in einen Wärmeofen innerhalb von 4 Stunden nach der vollständigen Verfestigung;
 Halten der Legierung in dem Wärmeofen bei einer ersten Temperatur von 600°F (316°C) bis 1800°F (982°C) über eine Dauer von mindestens 10 Stunden;
 Erhöhen der Ofentemperatur von der ersten Temperatur auf eine zweite Temperatur von mindestens 2125°F (1163°C) in einer solchen Weise, dass thermische Spannungen innerhalb der Legierung vermieden werden;
 Halten auf der zweiten Temperatur über eine Dauer von mindestens 10 Stunden;
 Vakuum-Lichtbogen-Umschmelzen einer VLU-Elektrode der Legierung mit einer Schmelzrate von 8 bis 11 lbs/Minute (3,63 bis 5 kg/Minute), um einen VLU-Block herzustellen.

2. Verfahren nach Anspruch 1, wobei die Legierung auf Nickelbasis Legierung 718 oder Legierung 706 ist.

3. Verfahren nach Anspruch 1 oder Anspruch 2, wobei das Gießen der Legierung auf Nickelbasis das Schmelzen und optionale Raffinieren der Legierung mittels Vakuuminduktionsschmelzen und/oder Argon-Sauerstoff-Entkohlung

und/oder Vakuum-Sauerstoff-Entkohlung umfasst.

4. Verfahren nach einem der vorangehenden Ansprüche, wobei das Glühen und Überaltern der Legierung das Erwärmen der Legierung bei mindestens 1200°F (649°C) über eine Dauer von mindestens 18 Stunden umfasst.

5. Verfahren nach einem der vorangehenden Ansprüche, wobei das Glühen und Überaltern der Legierung das Erwärmen der Legierung bei mindestens 1550°F (843°C) über eine Dauer von mindestens 10 Stunden umfasst.

6. Verfahren nach einem der vorangehenden Ansprüche, wobei das Elektroschlacke-Umschmelzen der Legierung das Elektroschlacke-Umschmelzen mit einer Schmelzrate von mindestens 10 lbs/Minute (4,54 kg/Minute) umfasst.

7. Verfahren nach einem der vorangehenden Ansprüche, wobei das Halten der Legierung in dem Wärmeofen das Halten der Legierung bei einer Ofentemperatur von mindestens 600°F (316°C) bis 1800°F (982°C) über eine Dauer von mindestens 20 Stunden umfasst.

8. Verfahren nach Anspruch 7, wobei das Halten der Legierung in dem Wärmeofen das Halten der Legierung bei einer Ofentemperatur von mindestens 900°F (482°C) bis 1800°F (982°C) über eine Dauer von mindestens 10 Stunden umfasst.

9. Verfahren nach einem der vorangehenden Ansprüche, wobei das Erhöhen der Ofentemperatur das Erhöhen der Ofentemperatur von der ersten Temperatur auf die zweite Temperatur in einem mehrstufigen Weise umfasst, umfassend:

Erhöhen der Ofentemperatur von der ersten Temperatur um nicht mehr als 100°F/Stunde (55,6°C/Stunde) auf eine Zwischentemperatur; und
weiteres Erhöhen der Ofentemperatur um nicht mehr als 200°F/Stunde (111°C/Stunde) von der Zwischentemperatur auf die zweite Temperatur.

10. Verfahren nach Anspruch 9, wobei die erste Temperatur niedriger als 1000°F (538°C) ist und die Zwischentemperatur mindestens 1000°F (538°C) beträgt.

11. Verfahren nach Anspruch 9, wobei die erste Temperatur niedriger als 1400°F (760°C) ist und die Zwischentemperatur mindestens 1400°F (760°C) beträgt.

12. Verfahren nach einem der vorangehenden Ansprüche, wobei die zweite Temperatur mindestens 2175°F (1191°C) beträgt.

13. Verfahren nach einem der vorangehenden Ansprüche, wobei die Legierung auf der zweiten Temperatur über eine Dauer von mindestens 24 Stunden gehalten wird.

14. Verfahren nach einem der vorangehenden Ansprüche, wobei das Elektroschlacke-Umschmelzen der Legierung einen ESU-Block mit einem Durchmesser erbringt, der größer als ein gewünschter Durchmesser der VLU-Elektrode ist, wobei das Verfahren im Anschluss an das Halten auf der zweiten Temperatur des Weiteren umfasst:

mechanische Bearbeitung des ESU-Blocks, um die Abmessungen des Blocks zu ändern und eine VLU-Elektrode mit dem gewünschten Durchmesser herzustellen.

15. Verfahren nach Anspruch 14, wobei der ESU-Block einen Durchmesser von ungefähr 34 Inch (864 mm) bis ungefähr 40 Inch (1016 mm) hat und die VLU-Elektrode einen kleineren Durchmesser von maximal ungefähr 34 Inch (864 mm) hat.

16. Verfahren nach Anspruch 14 oder 15, das im Anschluss an das Halten der Legierung auf der zweiten Temperatur und vor der mechanischen Bearbeitung des ESU-Blocks des Weiteren umfasst:

Abkühlen der Legierung auf eine Temperatur zur mechanischen Bearbeitung mit einer Abkühlrate von maximal 200°F/Stunde (111°C/Stunde).

17. Verfahren nach einem der vorangehenden Ansprüche, das im Anschluss an das Halten der Legierung auf der

zweiten Temperatur und vor dem Vakuum-Lichtbogen-Umschmelzen der VLU-Elektrode umfasst:

Abkühlen der Legierung von der zweiten Temperatur auf Raumtemperatur durch einen Abkühlprozess, der das Verringern der Ofentemperatur mit einer Rate von maximal 200°F/Stunde (111°C/Stunde) von der zweiten Temperatur auf eine erste Zwischentemperatur von maximal 1750°F (982°C) und das Halten auf der ersten Zwischentemperatur über eine Dauer von mindestens 10 Stunden umfasst.

18. Verfahren nach Anspruch 17, wobei das Abkühlen der Legierung des Weiteren umfasst:

Verringern der Ofentemperatur mit einer Rate von maximal 150°F/Stunde (83,3°C/Stunde) von der ersten Zwischentemperatur auf eine zweite Zwischentemperatur von maximal 1400°F (760°C) und Halten auf der zweiten Zwischentemperatur über eine Dauer von mindestens 5 Stunden.

19. Verfahren nach Anspruch 18, wobei die Legierung im Anschluss an das Halten auf der zweiten Zwischentemperatur in Luft auf ungefähr Raumtemperatur abgekühlt wird.

20. Verfahren nach einem der vorangehenden Ansprüche, das im Anschluss an das Halten der Legierung auf der zweiten Temperatur und vor der mechanischen Bearbeitung des ESU-Blocks Folgendes umfasst:

Abkühlen der Legierung von der zweiten Temperatur auf ungefähr Raumtemperatur in einer solchen Weise, dass thermische Spannungen in der Legierung vermieden werden; und Erwärmen der Legierung auf eine geeignete Temperatur zur mechanischen Bearbeitung in einer solchen Weise, dass thermische Spannungen in der Legierung vermieden werden.

21. Verfahren nach Anspruch 20, wobei das Erwärmen der Legierung auf eine geeignete Temperatur zur mechanischen Bearbeitung Folgendes umfasst:

Erwärmen der Legierung in einem Wärmeofen bei einer Ofentemperatur von mindestens 500°F (260°C) über eine Dauer von mindestens 2 Stunden;
Erhöhen der Ofentemperatur um mindestens ungefähr 20°F/Stunde (11,1°C/Stunde) auf mindestens 800°F (427°C);
weiteres Erhöhen der Ofentemperatur um mindestens ungefähr 30°F/Stunde (16,7°C/Stunde) auf mindestens 1200°F (649°C); und
weiteres Erhöhen der Ofentemperatur um mindestens ungefähr 40°F/Stunde (22,2°C/Stunde) auf eine Temperatur von mindestens 2025°F (1107°C) und Halten auf der Temperatur, bis die Legierung eine im Wesentlichen gleichmäßige Temperatur durch ihr gesamtes Volumen hindurch erreicht.

22. Verfahren nach einem der vorangehenden Ansprüche, wobei das Verfahren Folgendes umfasst:

Gießen einer Legierung auf Nickelbasis in einer Gießform, wobei die Superlegierung auf Nickelbasis Legierung 718 ist;
Glühen und Überaltern der Legierung durch Erwärmen der Legierung bei mindestens 1550°F (843°C) über eine Dauer von mindestens 10 Stunden;
Elektroschlack-Umschmelzen der Legierung mit einer Schmelzrate von mindestens 10 lbs/min (4,54 kg/min);
Verbringen der Legierung in einen Wärmeofen innerhalb von 4 Stunden nach der vollständigen Verfestigung nach dem Elektroschlack-Umschmelzen;
Halten der Legierung in dem Wärmeofen bei einer ersten Ofentemperatur von 900°F (482°C) bis 1800°F (982°C) über eine Dauer von mindestens 10 Stunden;
Erhöhen der Ofentemperatur um nicht mehr als 100°F/Stunde (55,6°C/Stunde) auf eine Ofenzwischentemperatur; und
weiteres Erhöhen der Ofentemperatur um nicht mehr als 200°F/Stunde (111°C/Stunde) von der Ofenzwischentemperatur auf eine zweite Ofentemperatur von mindestens 2125°F (1163°C) und Halten auf der zweiten Temperatur über eine Dauer von mindestens 10 Stunden; und
Vakuum-Lichtbogen-Umschmelzen einer VLU-Elektrode der Legierung mit einer Schmelzrate von 9 bis 10,25 lbs/Minute (4,09 bis 4,66 kg/min), um einen VLU-Block herzustellen.

23. Verfahren nach einem der vorangehenden Ansprüche, wobei der VLU-Block einen Durchmesser größer als 30 Inch (762 mm) hat.

24. Verfahren nach einem der vorangehenden Ansprüche, wobei der VLU-Block einen Durchmesser von mindestens 36 Inch (914 mm) hat.

25. Verfahren nach einem der vorangehenden Ansprüche, wobei das Gewicht des VLU-Blocks größer als 21.500 lbs (9772 kg) ist.

26. Verfahren nach einem der vorangehenden Ansprüche, wobei die Legierung auf Nickelbasis Folgendes umfasst:

50,0 bis 55,0 Gewichtsprozent Nickel;
17 bis 21,0 Gewichtsprozent Chrom;
0 bis 0,08 Gewichtsprozent Kohlenstoff;
0 bis 0,35 Gewichtsprozent Mangan;
0 bis 0,35 Gewichtsprozent Silicium;
2,8 bis 3,3 Gewichtsprozent Molybdän;
Niob und/oder Tantal, wobei die Summe aus Niob und Tantal 4,75 bis 5,5 Gewichtsprozent beträgt;
0,65 bis 1,15 Gewichtsprozent Titan;
0,20 bis 0,8 Gewichtsprozent Aluminium;
0 bis 0,006 Gewichtsprozent Bor; und
der Rest Eisen und beiläufige Verunreinigungen.

27. Verfahren nach Anspruch 26, wobei die Legierung auf Nickelbasis im Wesentlichen aus Folgendem besteht:

54,0 Gewichtsprozent Nickel;
0,5 Gewichtsprozent Aluminium;
0,01 Gewichtsprozent Kohlenstoff;
5,0 Gewichtsprozent Niob;
18 Gewichtsprozent Chrom;
3,0 Gewichtsprozent Molybdän;
0,9 Gewichtsprozent Titan; und
der Rest Eisen und beiläufige Verunreinigungen.

28. Verfahren nach Anspruch 22, wobei das Elektroschlack-Umschmelzen der Legierung einen ESU-Block mit einem Durchmesser erbringt, der größer als ein gewünschter Durchmesser der VLU-Elektrode ist, wobei das Verfahren des Weiteren umfasst:

Abkühlen der Legierung von der zweiten Temperatur auf eine geeignete Temperatur zur mechanischen Bearbeitung und dann mechanisches Bearbeiten der Legierung, um eine VLU-Elektrode mit dem gewünschten Durchmesser zu erhalten.

29. Verfahren nach Anspruch 22, wobei das Elektroschlack-Umschmelzen der Legierung einen ESU-Block mit einem Durchmesser erbringt, der größer als ein gewünschter Durchmesser der VLU-Elektrode ist, wobei das Verfahren des Weiteren Folgendes umfasst:

Abkühlen der Legierung von der zweiten Temperatur auf ungefähr Raumtemperatur in einer solchen Weise, dass thermische Spannungen in der Legierung vermieden werden;
Erwärmen der Legierung auf eine geeignete Temperatur zur mechanischen Bearbeitung in einer solchen Weise, dass thermische Spannungen in der Legierung vermieden werden;
mechanisches Bearbeiten der Legierung, um eine VLU-Elektrode mit dem gewünschten Durchmesser herzustellen.

Revendications

1. Procédé de production d'un superalliage à base de nickel qui est sensiblement dépourvu de ségrégations positives et négatives, le procédé comprenant :

le coulage d'un alliage qui est un superalliage à base de nickel dans un moule ;
le recuit et le surveillancement de l'alliage par chauffage de l'alliage à au moins 1 200 °F (649 °C) pendant au

moins 10 heures ;
la refusion sous laitier électroconducteur de l'alliage à une vitesse de fusion d'au moins 8 livres/min. (3,63 kg/min.) ;
le transfert de l'alliage vers un four de chauffage dans les 4 heures qui suivent la solidification totale ;
le maintien de l'alliage dans le four de chauffage à une première température de 600 °F (316 °C) à 1 800 °F (982 °C) pendant au moins 10 heures ;
l'augmentation de la température du four depuis la première température jusqu'à une deuxième température d'au moins 2 125 °F (1 163 °C) de manière à inhiber les contraintes thermiques dans l'alliage ;
le maintien à la deuxième température pendant au moins 10 heures ;
la refusion à l'arc sous vide d'une électrode refondue à l'arc sous vide (VAR) de l'alliage à une vitesse de fusion de 8 à 11 livres/minute (3,63 à 5 kg/minute) pour donner un lingot refondu à l'arc sous vide.

2. Procédé selon la revendication 1, dans lequel l'alliage à base de nickel est un parmi l'alliage 718 et l'alliage 706.

3. Procédé selon la revendication 1 ou la revendication 2, dans lequel le coulage de l'alliage à base de nickel comprend la fusion et éventuellement l'affinage de l'alliage par au moins un des procédés suivants : la fusion sous vide avec chauffage par induction, la décarburation à l'argon-oxygène et la décarburation sous vide à l'oxygène.

4. Procédé selon l'une quelconque des revendications précédentes, dans lequel le recuit et le survieillissement de l'alliage comprend le chauffage de l'alliage à au moins 1 200 °F (649 °C) pendant au moins 18 heures.

5. Procédé selon l'une quelconque des revendications précédentes, dans lequel le recuit et le survieillissement de l'alliage comprend le chauffage de l'alliage à au moins 1 550 °F (843 °C) pendant au moins 10 heures.

6. Procédé selon l'une quelconque des revendications précédentes, dans lequel la refusion sous laitier électroconducteur de l'alliage comprend la refusion sous laitier électroconducteur à une vitesse de fusion d'au moins 10 livres/minute (4,54 kg/minute).

7. Procédé selon l'une quelconque des revendications précédentes, dans lequel le maintien de l'alliage dans le four de chauffage comprend le maintien de l'alliage à une température du four d'au moins 600 °F (316 °C) jusqu'à 1 800 °F (982 °C) pendant au moins 20 heures.

8. Procédé selon la revendication 7, dans lequel le maintien de l'alliage dans le four de chauffage comprend le maintien de l'alliage à une température du four d'au moins 900 °F (482 °C) jusqu'à 1 800 °F (982 °C) pendant au moins 10 heures.

9. Procédé selon l'une quelconque des revendications précédentes, dans lequel l'augmentation de la température du four comprend l'augmentation de la température du four depuis une première température jusqu'à une deuxième température en plusieurs étapes comprenant :

l'augmentation de la température du four depuis la première température par pas plus de 100 °F/heure (55,6 °C/heure) jusqu'à une température intermédiaire ; et

l'augmentation supplémentaire de la température du four par pas plus de 200 °F/heure (111 °C/heure) depuis la température intermédiaire jusqu'à la deuxième température.

10. Procédé selon la revendication 9, dans lequel la première température est inférieure à 1 000 °F (538 °C) et la température intermédiaire est d'au moins 1 000 °F (538 °C).

11. Procédé selon la revendication 9, dans lequel la première température est inférieure à 1 400 °F (760 °C) et la température intermédiaire est d'au moins 1 400 °F (760 °C).

12. Procédé selon l'une quelconque des revendications précédentes, dans lequel la deuxième température est d'au moins 2 175 °F (1 191 °C).

13. Procédé selon l'une quelconque des revendications précédentes, dans lequel l'alliage est maintenu à la deuxième température pendant au moins 24 heures.

14. Procédé selon l'une quelconque des revendications précédentes, dans lequel la refusion sous laitier électrocon-

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ducteur de l'alliage donne un lingot refondu sous laitier électroconducteur ayant un diamètre supérieur au diamètre désiré de l'électrode refondue à l'arc sous vide, le procédé comprenant en outre, après le maintien à la deuxième température:

5 le travail mécanique du lingot refondu sous laitier électroconducteur pour faire varier les dimensions du lingot et pour donner une électrode refondue à l'arc sous vide ayant le diamètre désiré.

15. Procédé selon la revendication 14, dans lequel le lingot refondu sous laitier électroconducteur a un diamètre d'environ 34 pouces (864 mm) à environ 40 pouces (1 016 mm) et l'électrode refondue à l'arc sous vide a un diamètre plus petit, de pas plus d'environ 34 pouces (864 mm).

16. Procédé selon la revendication 14 ou 15, comprenant en outre, après le maintien de l'alliage à la deuxième température et avant le travail mécanique du lingot refondu sous laitier électroconducteur :

15 le refroidissement de l'alliage jusqu'à une température de travail mécanique à une vitesse de refroidissement de pas plus de 200 °F/heure (111 °C/heure).

17. Procédé selon l'une quelconque des revendications précédentes, comprenant en outre, après le maintien de l'alliage à la deuxième température et avant la refusion à l'arc sous vide de l'électrode refondue à l'arc sous vide :

20 le refroidissement de l'alliage depuis la deuxième température jusqu'à la température ambiante par un procédé de refroidissement comprenant la réduction de la température du four à une vitesse de pas plus de 200 °F/heure (111 °C/heure) depuis la deuxième température jusqu'à une première température intermédiaire de pas plus de 1 750 °F (982 °C) et le maintien à la première température intermédiaire pendant au moins 10 heures.

25 18. Procédé selon la revendication 17, dans lequel le refroidissement de l'alliage comprend en outre :

la réduction de la température du four à une vitesse de pas plus de 150 °F/heure (83,3 °C/heure) depuis la première température intermédiaire jusqu'à une deuxième température intermédiaire de pas plus de 1 400 °F (760 °C) et le maintien à la deuxième température intermédiaire pendant au moins 5 heures.

19. Procédé selon la revendication 18, dans lequel après le maintien à la deuxième température intermédiaire, l'alliage est refroidi à l'air jusqu'à environ la température ambiante.

35 20. Procédé selon l'une quelconque des revendications précédentes, comprenant en outre, après le maintien de l'alliage à la deuxième température et avant le travail mécanique du lingot refondu sous laitier électroconducteur :

le refroidissement de l'alliage depuis la deuxième température jusqu'à environ la température ambiante de manière à inhiber les contraintes thermiques dans l'alliage ; et

40 le chauffage de l'alliage jusqu'à une température de travail mécanique appropriée d'une manière qui inhibe les contraintes thermiques dans l'alliage.

21. Procédé selon la revendication 20, dans lequel le chauffage de l'alliage jusqu'à une température de travail mécanique appropriée comprend :

45 le chauffage de l'alliage dans un four de chauffage à une température du four d'au moins 500 °F (260 °C) pendant au moins 2 heures ;

l'augmentation de la température du four par au moins environ 20 °F/heure (11,1 °C/heure) jusqu'à au moins 800 °F (427 °C) ;

50 l'augmentation supplémentaire de la température du four par au moins environ 30 °F/heure (16,7 °C/heure) jusqu'à au moins 1 200 °F (649 °C) ; et

l'augmentation supplémentaire de la température du four par au moins environ 40 °F/heure (22,2 °C/heure) jusqu'à une température d'au moins 2 025 °F (1 107 °C) et le maintien à cette température jusqu'à ce que l'alliage ait partout une température sensiblement uniforme.

55 22. Procédé selon l'une quelconque des revendications précédentes, le procédé comprenant :

le coulage d'un alliage à base de nickel dans un moule, dans lequel le superalliage à base de nickel est l'alliage

718 ;

le recuit et le surveillissement de l'alliage par chauffage de l'alliage à au moins 1 550 °F (843 °C) pendant au moins 10 heures ;

la refusion sous laitier électroconducteur de l'alliage à une vitesse de fusion d'au moins 10 livres/min. (4,54 kg/min.) ;

le transfert de l'alliage vers un four de chauffage dans les 4 heures qui suivent la solidification totale après refusion sous laitier électroconducteur ;

le maintien de l'alliage dans le four de chauffage à une première température du four de 900 °F (482 °C) à 1 800 °F (982 °C) pendant au moins 10 heures ;

l'augmentation de la température du four par pas plus 100 °F/heure (55,6 °C/heure) jusqu'à une température du four intermédiaire ;

l'augmentation supplémentaire de la température du four par pas plus de 200 °F/heure (111 °C/heure) depuis la température du four intermédiaire jusqu'à une deuxième température du four d'au moins 2 125 °F (1 163 °C), et le maintien à la deuxième température pendant au moins 10 heures; et

la refusion à l'arc sous vide d'une électrode refondue à l'arc sous vide (VAR) de l'alliage à une vitesse de fusion de 9 à 10,25 livres/minute (4,09 à 4,66 kg/minute) pour donner un lingot refondu à l'arc sous vide.

23. Procédé selon l'une quelconque des revendications précédentes, dans lequel le lingot refondu à l'arc sous vide a un diamètre supérieur à 30 pouces (762 mm).

24. Procédé selon l'une quelconque des revendications précédentes, dans lequel le lingot refondu à l'arc sous vide a un diamètre d'au moins 36 pouces (914 mm).

25. Procédé selon l'une quelconque des revendications précédentes, dans lequel le poids du lingot refondu à l'arc sous vide est supérieur à 21 500 livres (9 772 kg).

26. Procédé selon l'une quelconque des revendications précédentes, dans lequel l'alliage à base de nickel comprend :

50,0 à 55,0 pour cent en poids de nickel;

17 à 21,0 pour cent en poids de chrome ;

0 jusqu'à 0,08 pour cent en poids de carbone ;

0 jusqu'à 0,35 pour cent en poids de manganèse ;

0 jusqu'à 0,35 pour cent en poids de silicium ;

2,8 jusqu'à 3,3 pour cent en poids de molybdène ;

au moins un parmi le niobium et le tantale dans lequel la somme du niobium et du tantale est de

4,75 jusqu'à 5,5 pour cent en poids ;

0,65 jusqu'à 1,15 pour cent en poids de titane ;

0,20 jusqu'à 0,8 pour cent en poids d'aluminium ;

0 jusqu'à 0,006 pour cent en poids de bore ; et

le reste étant du fer et des impuretés accidentelles.

27. Procédé selon la revendication 26, dans lequel l'alliage à base de nickel est constitué essentiellement de :

54,0 pour cent en poids de nickel ;

0,5 pour cent en poids d'aluminium ;

0,01 pour cent en poids de carbone ;

5,0 pour cent en poids de niobium ;

18,0 pour cent en poids de chrome ;

3,0 pour cent en poids de molybdène ;

0,9 pour cent en poids de titane ; et

le reste étant du fer et des impuretés accidentelles.

28. Procédé selon la revendication 22, dans lequel la refusion sous laitier électroconducteur de l'alliage donne un lingot refondu sous laitier électroconducteur ayant un diamètre supérieur à un diamètre désiré de l'électrode refondue à l'arc sous vide, le procédé comprenant en outre :

le refroidissement de l'alliage depuis la deuxième température jusqu'à une température de travail mécanique appropriée et ensuite le travail mécanique de l'alliage pour donner une électrode refondue à l'arc sous vide

ayant le diamètre désiré.

29. Procédé selon la revendication 22, dans lequel la refusion sous laitier électroconducteur de l'alliage donne un lingot refondu sous laitier électroconducteur ayant un diamètre supérieur à un diamètre désiré de l'électrode refondue à l'arc sous vide, le procédé comprenant en outre:

le refroidissement de l'alliage depuis la deuxième température jusqu'à environ la température ambiante d'une manière qui inhibe les contraintes thermiques dans l'alliage;

le chauffage de l'alliage jusqu'à une température de travail mécanique appropriée d'une manière qui inhibe les contraintes thermiques dans l'alliage ; et

le travail mécanique de l'alliage pour donner une électrode refondue à l'arc sous vide ayant le diamètre désiré.

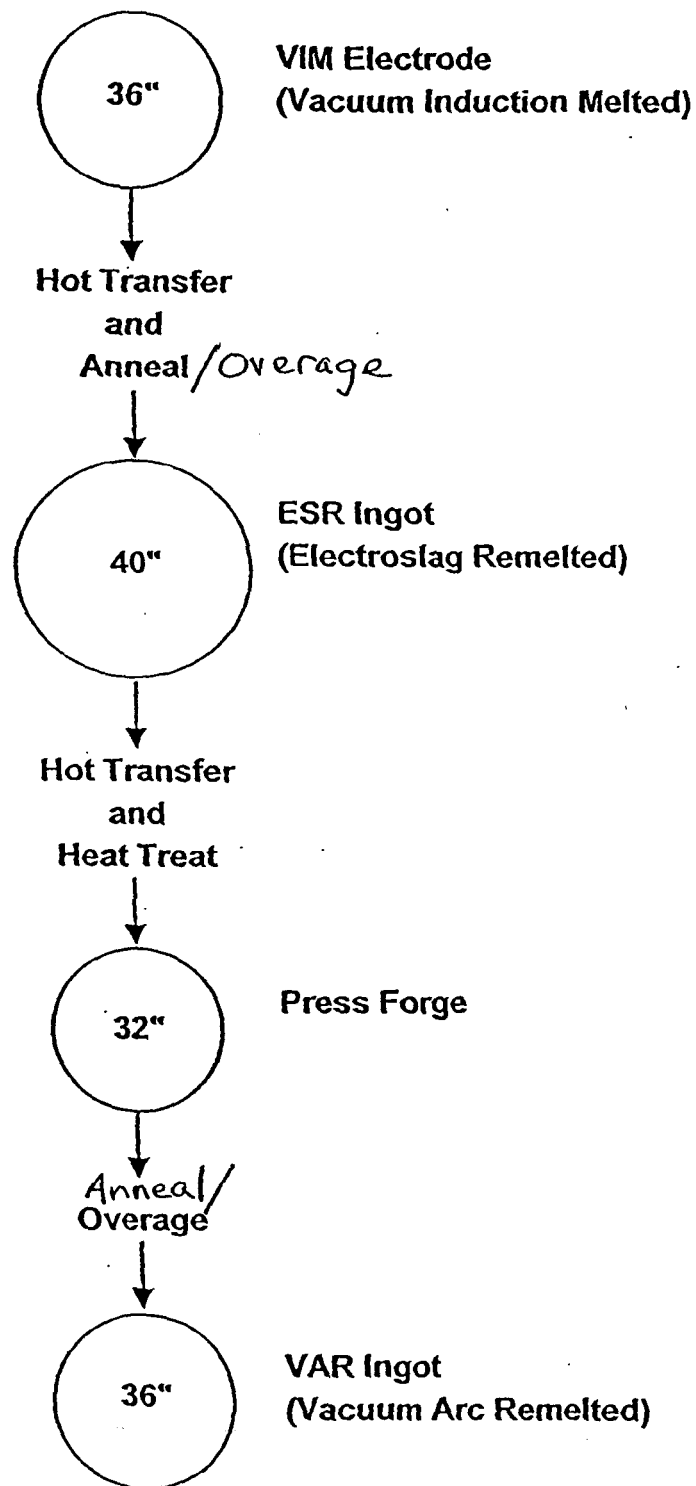


FIGURE 1

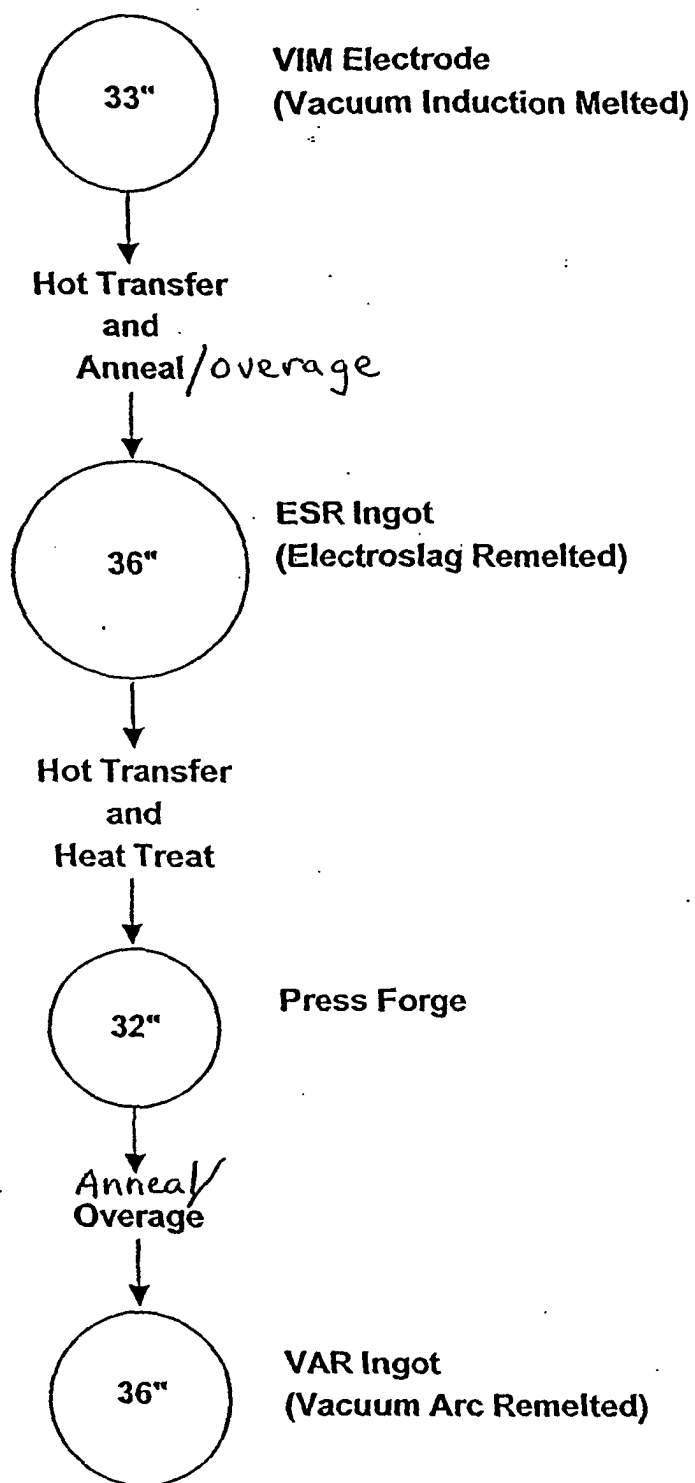


FIGURE 2

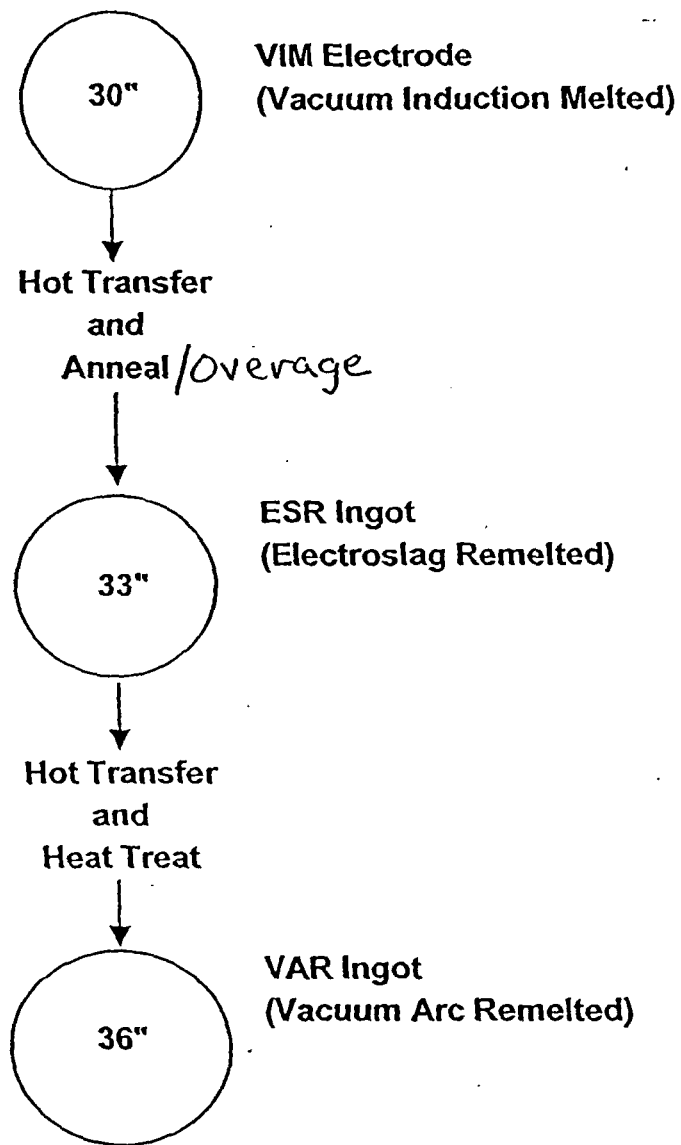


FIGURE 3

REFERENCES CITED IN THE DESCRIPTION

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Non-patent literature cited in the description

- **MOYER et al.** *The Minerals, Metals and Materials Society*, 1994, 39-48 [0008]